Commission Decision of 21 February 2008 on additional guarantees in intra-Community trade of pigs relating to Aujeszky's disease and criteria to provide information on this disease (notified under document number C(2008) 669) (Codified version) (Text with EEA relevance) (2008/185/EC)

COMMISSION DECISION

of 21 February 2008

on additional guarantees in intra-Community trade of pigs relating to Aujeszky's disease and criteria to provide information on this disease

(notified under document number C(2008) 669)

(Codified version)

(Text with EEA relevance)

(2008/185/EC)

THE COMMISSION OF THE EUROPEAN COMMUNITIES,

Having regard to the Treaty establishing the European Community,

Having regard to Council Directive 64/432/EEC of 26 June 1964 on animal health problems affecting intra-Community trade in bovine animals and swine[(1)], and in particular Article 8, Article 9(2) and Article 10(2) thereof,

Whereas:

(1) Commission Decision 2001/618/EC of 23 July 2001 on additional guarantees in intra-Community trade of pigs relating to Aujeszky's disease, criteria to provide information on this disease and repealing Decisions 93/24/EEC and 93/244/EEC[(2)] has been substantially amended several times[(3)]. In the interests of clarity and rationality the said Decision should be codified.

(2) The International Office of Epizootic Diseases (OIE) is the international organisation designated under the Agreement on the Application of Sanitary and Phytosanitary Measures in application of GATT 1994 which is responsible for the establishment of international animal health rules for trade in animals and animal products. These rules are published in the International Animal Health Code.

(3) The chapter of the International Animal Health Code on Aujeszky's disease has been substantially amended.

(4) It is appropriate to modify the additional guarantees required in intra-Community trade of pigs in relation to Aujeszky's disease in order to ensure their consistency with the international rules on this disease and better control in the Community.

(5) Criteria must be established on the information to be provided by the Member States on Aujeszky's disease, in accordance with Article 8 of Directive 64/432/EEC.

(6) The measures provided for in this Decision are in accordance with the opinion of the Standing Committee on the Food Chain and Animal Health,

HAS ADOPTED THIS DECISION:

Article 1 U.K.

[[F1]Great Britain is free of Aujeszky's disease and vaccination is prohibited in Great Britain. The member States or regions of such States that are recognised as free from the disease are listed in Annex I, and those that are not free but have EU approved Aujeszky's disease eradication plans in place are listed in Annex II. Pigs intended for breeding or production dispatched to Great Britain must come from a member State or region thereof listed in Annex I or must comply with the following additional conditions:**]**

1. Aujeszky's disease must be compulsorily notifiable in the Member State of origin;

2. a plan for the control and eradication of Aujeszky's disease, fulfilling the criteria laid down in Article 9(1) of Directive 64/432/EEC, must be in place in the Member State or regions of origin under the supervision of the competent authority. Appropriate measures on pig transport and movements must be in place according to this plan for preventing a spread of disease between holdings of a different status;

3. with regard to the holding of origin of the pigs:

   (a) no clinical, pathological or serological evidence of Aujeszky's disease has been recorded in the previous 12 months in the holding in question;

   (b) no clinical, pathological or serological evidence of Aujeszky's disease has been recorded in the previous 12 months in the holdings located in an area of 5 km surrounding the holding of origin of the pigs; however, this provision shall not apply if, in these latter holdings, disease monitoring and eradication measures have been regularly applied under the supervision of the competent authority and in accordance with the eradication plan referred to in point (2), and these measures have effectively prevented any spread of disease to the holding in question;

   (c) vaccination against Aujeszky's disease has not been carried out for at least 12 months;

   (d) the pigs have been subjected on at least two occasions at an interval of at least four months to a serological survey for the presence of ADV-gE or ADV-gB or ADV-gD antibody or to the whole Aujeszky's disease virus. This survey must have shown the absence of Aujeszky's disease and that vaccinated pigs have been free from gE antibodies;

   (e) no pigs have been introduced from holdings of a lower animal health status as regards Aujeszky's disease in the previous 12 months, unless they have been tested for Aujeszky's disease with negative results;

4. the pigs to be moved:

   (a) have not been vaccinated;

(b) have been kept isolated in accommodation approved by the competent authority, during the 30 days prior to movement, and in such a way that any risk of spreading Aujeszky's disease to these pigs is prevented;

(c) must have lived in the holding of origin or in a holding of an equivalent status since birth, and have remained in the holding of origin for at least:

(i) 30 days, in the case of pigs intended for production;

(ii) 90 days, in the case of pigs intended for breeding;

(d) have been subjected with negative results to at least two serological tests for ADV-gB or ADV-gD or the whole Aujeszky's disease virus, at a distance of at least 30 days between each test. However, in case of pigs less than four months old, the serological test for ADV-gE may also be used. Sampling for the last test must be performed within 15 days prior to shipment. The number of pigs tested in the isolation unit must be sufficient to detect:

(i) 2 % seroprevalence with 95 % confidence in the isolation unit in case of pigs intended for production;

(ii) 0,1 % seroprevalence with 95 % confidence in the isolation unit in case of pigs intended for breeding.

However, the first of the two tests shall not be necessary if:

(i) in the framework of the plan referred to in point (2), a serological survey has been carried out in the holding of origin between 45 and 170 days prior to shipment, demonstrating the absence of Aujeszky's disease antibodies and that vaccinated pigs have been free from gE antibodies;

(ii) the pigs to be moved have lived in the holding of origin since birth;

(iii) no pigs have moved on to the holding of origin while the pigs to be moved have been kept in isolation.

Textual Amendments

F1 Words in Art. 1 substituted (31.12.2020) by The Import of, and Trade in, Animals and Animal Products (Miscellaneous Amendments) (EU Exit) Regulations 2020 (S.I. 2020/1462), regs. 1(3), **51(2)** (with regs. 69-71)

Article 2 *U.K.*

[[F2]Pigs intended for slaughter dispatched to Great Britain must come from a member State or region thereof listed in Annex I or must comply with the following additional conditions:**]**

1. Aujeszky's disease must be compulsorily notifiable in the Member State of origin;

2. a plan for the control and eradication of Aujeszky's disease is in place in the Member State or regions of origin of the pigs, fulfilling the criteria laid down in Article 1(2);

3. all the pigs in question must be transported directly to the slaughterhouse of destination and either:

(a) they come from a holding which fulfils the conditions laid down in Article 1(3); or

(b) they have been vaccinated against Aujeszky's disease at least 15 days prior to their shipment and come from a holding of origin where:

(i) in the framework of the plan referred to in point (2), Aujeszky's disease monitoring and eradication measures have been regularly applied under the supervision of the competent authority for the previous 12 months;

(ii) they had remained for at least 30 days before dispatch and where no clinical or pathological evidence of this disease has been detected at the moment of completion of the health certificate **[**[F3]accompanying the consignment**]**; or

(c) they have not been vaccinated and they proceed from a holding where:

(i) in the framework of the plan referred to in point 2, Aujeszky's disease monitoring and eradication measures have been regularly applied under the supervision of the competent authority in the previous 12 months and no clinical, pathological or serological evidence of Aujeszky's disease has been recorded in the previous six months;

(ii) vaccination against Aujeszky's disease and introduction of vaccinated pigs have been forbidden by the competent authority, since the holding is in the process of reaching the highest status as regards Aujeszky's disease in accordance to the plan referred to in point (2);

(iii) they have lived for at least 90 days before dispatch.

Textual Amendments

F2 Words in Art. 2 substituted (31.12.2020) by The Import of, and Trade in, Animals and Animal Products (Miscellaneous Amendments) (EU Exit) Regulations 2020 (S.I. 2020/1462), regs. 1(3), **51(3)(a)** (with regs. 69-71)

F3 Words in Art. 2(3)(b)(ii) substituted (31.12.2020) by The Import of, and Trade in, Animals and Animal Products (Miscellaneous Amendments) (EU Exit) Regulations 2020 (S.I. 2020/1462), regs. 1(3), **51(3)(b)** (with regs. 69-71)

[F4]Article 3 U.K.

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Textual Amendments

F4 Art. 3 omitted (31.12.2020) by virtue of The Import of, and Trade in, Animals and Animal Products (Miscellaneous Amendments) (EU Exit) Regulations 2020 (S.I. 2020/1462), regs. 1(3), **51(4)** (with regs. 69-71)

F5 Article 4 U.K.

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Textual Amendments

F5 Art. 4 omitted (31.12.2020) by virtue of The Import of, and Trade in, Animals and Animal Products (Miscellaneous Amendments) (EU Exit) Regulations 2020 (S.I. 2020/1462), regs. 1(3), **51(4)** (with regs. 69-71)

Article 5 U.K.

The serological tests carried out to monitor or detect Aujeszky's disease in pigs in accordance with this Decision must meet the standards laid down in Annex III.

F6 Article 6 U.K.

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Textual Amendments

F6 Art. 6 omitted (31.12.2020) by virtue of The Import of, and Trade in, Animals and Animal Products (Miscellaneous Amendments) (EU Exit) Regulations 2020 (S.I. 2020/1462), regs. 1(3), **51(5)** (with regs. 69-71)

[F7 Article 7 U.K.

In the case of pigs destined for Great Britain from a member State, it is a requirement that the official veterinarian in the country of origin has ascertained the disease status of the holding of origin and the disease-free status of the region of origin and checked the compliance of the pigs in question with the conditions laid down in this Decision.]

Textual Amendments

F7 Art. 7 substituted (31.12.2020) by The Import of, and Trade in, Animals and Animal Products (Miscellaneous Amendments) (EU Exit) Regulations 2020 (S.I. 2020/1462), regs. 1(3), **51(6)** (with regs. 69-71)

[F8 Article 8 U.K.

When pigs are transported to Great Britain, it is a requirement that the exporting member State has ensured that the pigs do not come into contact with pigs of different or unknown status, as regards Aujeszky's disease, during transport or transit.]

Textual Amendments

F8 Art. 8 substituted (31.12.2020) by The Import of, and Trade in, Animals and Animal Products (Miscellaneous Amendments) (EU Exit) Regulations 2020 (S.I. 2020/1462), regs. 1(3), **51(7)** (with regs. 69-71)

Article 9 U.K.

Decision 2001/618/EC is repealed.

References to the repealed Decision shall be construed as references to this Decision and shall be read in accordance with the correlation table in Annex VI.

F9 Article 10 U.K.

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Textual Amendments

F9 Art. 10 omitted (31.12.2020) by virtue of The Import of, and Trade in, Animals and Animal Products (Miscellaneous Amendments) (EU Exit) Regulations 2020 (S.I. 2020/1462), regs. 1(3), **51(8)** (with regs. 69-71)

[F10ANNEX I U.K.

Member States or regions thereof free of Aujeszky's disease and where vaccination is prohibited

Textual Amendments

F10 Substituted by Commission Implementing Decision (EU) 2019/1970 of 26 November 2019 amending Annex II to Decision 93/52/EEC as regards the officially brucellosis (B. melitensis)-free status and Annex II to Decision 2003/467/EC as regards the officially brucellosis-free status of certain regions of Spain and Annexes I and II to Decision 2008/185/EC as regards the free status and the approval of the eradication programmes for Aujeszky's disease for certain regions of Italy (notified under document C(2019) 8378) (Text with EEA relevance).

ISO code	Member State	Regions
BE	Belgium	All regions
CZ	Czechia	All regions
DK	Denmark	All regions
DE	Germany	All regions
IE	Ireland	All regions
FR	France	The departments of Ain, Aisne, Allier, Alpes-de-Haute-Provence, Alpes-Maritimes, Ardèche, Ardennes, Ariège, Aube, Aude, Aveyron, Bas-Rhin, Bouches-du-Rhône, Calvados, Cantal, Charente, Charente-Maritime, Cher, Corrèze, Côte-d'Or, Côtes-d'Armor, Creuse, Deux-Sèvres, Dordogne, Doubs, Drôme, Essonne, Eure, Eure-et-Loir, Finistère, Gard, Gers, Gironde, Hautes-Alpes, Hauts-de-Seine, Haute Garonne, Haute-Loire, Haute-Marne, Hautes-Pyrénées, Haut-Rhin, Haute-Saône, Haute-Savoie, Haute-Vienne, Hérault, Indre, Ille-et-Vilaine, Indre-et-Loire, Isère, Jura, Landes, Loire, Loire-Atlantique, Loir-et-Cher, Loiret, Lot, Lot-et-Garonne, Lozère, Maine-et-Loire, Manche, Marne, Mayenne, Meurthe-et-Moselle, Meuse, Morbihan,

		Moselle, Nièvre, Nord, Oise, Orne, Paris, Pas-de-Calais, Pyrénées-Atlantiques, Pyrénées-Orientales, Puy-de-Dôme, Réunion, Rhône, Sarthe, Saône-et-Loire, Savoie, Seine-et-Marne, Seine-Maritime, Seine-Saint-Denis, Somme, Tarn, Tarn-et-Garonne, Territoire de Belfort, Val-de-Marne, Val-d'Oise, Var, Vaucluse, Vendée, Vienne, Vosges, Yonne, Yvelines
IT	Italy	Autonomous Province of Bolzano Region Friuli Venezia Giulia
CY	Cyprus	All regions
LU	Luxembourg	All regions
HU	Hungary	All regions
NL	Netherlands	All regions
AT	Austria	All regions
PL	Poland	Voivodship podlaskie the following powiaty: augustowski, białostocki, Białystok, bielski, hajnowski, moniecki, sejneński, siemiatycki, sokólski, suwalski, Suwałki
SI	Slovenia	All regions
SK	Slovakia	All regions
FI	Finland	All regions
SE	Sweden	All regions
F11 . . .	**F11** . . .	**F11** . . .

Textual Amendments

F11 Words in Annex 1 omitted (31.12.2020) by virtue of The Import of, and Trade in, Animals and Animal Products (Miscellaneous Amendments) (EU Exit) Regulations 2020 (S.I. 2020/1462), regs. 1(3), **51(9)** (with regs. 69-71)

ANNEX II U.K.

Member States or regions thereof where approved national control programmes for the eradication of Aujeszky's disease are in place

ISO code	Member State	Regions
ES	Spain	All regions
IT	Italy	Region Emilia-Romagna Region Lombardia Region Piemonte Region Umbria Region Veneto
LT	Lithuania	All regions
PL	Poland	Voivodship dolnośląskie: all powiaty; Voivodship kujawsko-pomorskie: all powiaty; Voivodship lubelskie: all powiaty; Voidodship lubuskie: all powiaty; Voivodship łódzkie: all powiaty; Voivodship małopolskie: all powiaty; Voivodship mazowieckie: all powiaty; Voivodship opolskie: all powiaty; Voivodship podkarpackie: all powiaty; Voivodship podlaskie the following powiaty: grajewski, kolneński, łomżyński, Łomża, wysokomazowiecki, zambrowski. Voivodship pomorskie: all powiaty; Voivodship śląskie: all powiaty; Voivodship świętokrzyskie: all powiaty; Voivodship warmińsko-mazurskie: all powiaty; Voivodship wielkopolskie: all powiaty; Voivodship zachodniopomorskie: all powiaty.]

ANNEX III U.K.

Standards for Aujeszky's disease serological tests — Protocol for the enzyme linked immunosorbent assay (ELISA) for detecting antibodies to Aujeszky's disease virus (whole virus), to glycoprotein B (ADV-gB), to glycoprotein D (ADV-gD) or to glycoprotein E (ADV-gE)

1. The institutes listed in paragraph 2(d) shall evaluate Elisa ADV-gE tests and kits against the criteria in paragraph 2(a), (b) and (c). The competent authority in each Member State shall ensure that only Elisa ADV-gE kits that meet these standards shall be registered. The examinations listed in 2(a) and (b) must be carried out prior to approval of the test and the examination in 2(c), at least, must thereafter be carried out on each batch.

2. Standardisation, sensitivity and specificity of the test. U.K.

(a) The sensitivity of the test must be of such a level that the following Community reference sera are scored positive:

— Community reference serum ADV 1 at 1:8 dilution,
— Community reference serum ADV-gE A,
— Community reference serum ADV-gE B,
— Community reference serum ADV-gE C,
— Community reference serum ADV-gE D,
— Community reference serum ADV-gE E,
— Community reference serum ADV-gE F.

(b) The specificity of the test must be of such a level that the following Community reference sera are scored negative:

— Community reference serum ADV-gE G,
— Community reference serum ADV-gE H,
— Community reference serum ADV-gE J,
— Community reference serum ADV-gE K,
— Community reference serum ADV-gE L,
— Community reference serum ADV-gE M,
— Community reference serum ADV-gE N,
— Community reference serum ADV-gE O,
— Community reference serum ADV-gE P,
— Community reference serum ADV-gE Q.

(c) For batch control, Community reference serum ADV 1 must be scored positive at 1:8 dilution and one of the Community reference sera from ADV-gE G to ADV-gE Q, as listed in point (b), must be scored negative.

For batch control of ADV-gB and ADV-gD kits, Community reference serum ADV 1 must be scored positive at the dilution of 1:2 and Community reference serum Q referred to in (b) should be scored negative.

(d) [F12The institutes listed below will, in addition, be responsible for checking the quality of the ELISA method in each Member State, and in particular for producing and standardising national reference sera according to the Community reference sera.

AT	AGES: Österreichische Agentur für Gesundheit und Ernährungssicherheit GmbH — Institut für veterinärmedizinische Untersuchungen Mödling (Austrian Agency for Health and Consumer Protection — Institute for veterinary investigations Mödling) Robert Koch-Gasse 17 A-2340 Mödling Tel. +43 (0) 505 55-38112 Fax +43 (0) 505 55-38108 Email: vetmed.moedling@ages.at
BE	CODA — CERVA — VAR Veterinary and Agrochemical Research Centre Groeselenberg 99 B-1180 Brussels
CY	State Veterinary Laboratory Veterinary Services 1417 Athalassa Nicosia
CZ	Státní veterinární ústav Olomouc Jakoubka ze Stříbra 1 779 00 Olomouc Telefon: 585 557 111 Fax 585 222 394 email: svuolomouc@svuol.cz
DE	Friedrich-Loeffler-Institut Bundesforschungsinstitut für Tiergesundheit Südufer 10 D-17493 Greifswald — Insel Riems Tel. + 49 38351 7-0 Fax + 49 38351 7-1219, 7-1151, 7-1226
DK	National Veterinary Institute Technical University of Denmark Lindholm Island DK-4774 Kalvehave Denmark Switchboard: +45 88 60 00 Fax +45 88 79 01 Email: vet@vet.dtu.dk
EE	Veterinaar- ja Toidulaboratoorium Kreutzwaldi 30, 51006 Tartu, Estonia Tel. + 372 7 386 100

	Faks: + 372 7 386 102 Email: info@vetlab.ee
ES	Laboratorio Central de Sanidad Animal de Algete Carretera de Algete, km 8 Algete 28110 (Madrid) Tel. +34 916 290 300 Fax +34 916 290 598 Email: lcv@mapya.es
FI	Finnish Food Safety Authority Animal Diseases and Food Safety Research Mustialankatu 3 FI-00790 Helsinki, Finland Email: info@evira.fi Tel. +358 20 772 003 (exchange) Fax +358 20 772 4350
FR	Laboratoire d'études et de recherches avicoles, porcines et piscicoles AFSSA site de Ploufragan/Brest — LERAPP BP 53 22440 Ploufragan
F13 . . .	**F13** . . .
GR	Centre of Athens Veterinary Institutes 25 Neapoleos Street, GR-153 10 Agia Paraskevi Attiki Tel. +30 2106010903
HU	Nemzeti Élelmiszerlánc-biztonsági Hivatal, Állat-egészségügyi Diagnosztikai Igazgatóság Central Agricultural Office, Veterinary Diagnostic Directorate Address: 1149 Budapest, Tábornok u. 2. Mailing Address: 1581 Budapest, 146. Pf. 2. Tel. +36 1 460-6300 Fax +36 1 252-5177 Email: ugyfelszolgalat@nebih.gov.hu
IE	Virology Division Central Veterinary Research Laboratory Department of Agriculture and Food Laboratories Backweston Campus Stacumny Lane Celbridge Co. Kildare

IT	Centro di referenza nazionale per la malattia di Aujeszky — Pseudorabbia c/o Istituto zooprofilattico sperimentale della Lombardia e dell'Emilia Romagna, Via Bianchi, 9; 25124 Brescia
LT	National Veterinary Laboratory (Nacionalinė veterinarijos laboratorija) J. Kairiūkščio 10 LT-08409 Vilnius
LU	CODA — CERVA — VAR Veterinary and Agrochemical Research Centre Groeselenberg 99 B-1180 Brussels
LV	Pārtikas drošības, dzīvnieku veselības un vides zinātniskais institūts 'BIOR' (Institute of Food Safety, Animal Health and Environment BIOR) Lejupes iela 3, Rīga, LV-1076 Tel. +371 76205 13 Fax +371 7620434 Email: bior@bior.lv
MT	National Veterinary Laboratory Veterinary and Phytosanitary Regulation Department Ministry for Sustainable Development, the Environment and Climate Change, Abattior Square, Albert Town, Triq Prince Albert, Marsa, Malta Tel. +356 22925389
NL	Centraal Instituut voor Dierziekte Controle CIDC-Lelystad Hoofdvestiging: Houtribweg 39 Nevenvestiging: Edelhertweg 15 Postbus 2004 8203 AA Lelystad
PL	Laboratory Department of Swine Diseases Państwowy Instytut Weterynaryjny — Państwowy Instytut Badawczy al. Partyzantów 57, 24-100 Puławy Tel. +48 81 889 30 00 Fax +48 81 886 25 95

	Email: sekretariat@piwet.pulawy.pl
PT	Laboratório Nacional de Investigação Veterinária (LNIV) Estrada de Benfica, 701 P-1549-011 Lisboa
RO	Laboratorul Național de Referință pentru Herpesviroze Institutul de Diagnostic și Sănătate Animală Str. Dr Staicovici, nr. 6, cod 050557, sector 5, București telefon: 0374.322.015 fax 0214.113.394 email: office@idah.ro
SE	Statens veterinärmedicinska anstalt Department of Virology S-751 89 Uppsala Tel. (46-18) 67 40 00 Fax (46-18) 67 44 67
SI	Univerza v Ljubljani Veterinarska fakulteta Nacionalni veterinarski inštitut Gerbičeva 60, SI-1000 Ljubljana
SK	Štátny veterinárny ústav Pod dráhami 918 960 86 Zvolen Slovenska republika]

Textual Amendments

F12 Substituted by Commission Implementing Decision (EU) 2016/1782 of 5 October 2016 amending Decision 2008/185/EC as regards the inclusion of Lithuania in the list of Member States where an approved national control programme for Aujeszky's disease is in place and updating the list of national institutes in Annex III (notified under document C(2016) 6288) (Text with EEA relevance).

F13 Words in Annex 3 omitted (31.12.2020) by virtue of The Import of, and Trade in, Animals and Animal Products (Miscellaneous Amendments) (EU Exit) Regulations 2020 (S.I. 2020/1462), regs. 1(3), **51(10)** (with regs. 69-71)

[F14]ANNEX IV U.K.

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Textual Amendments

F14 Annex 4 omitted (31.12.2020) by virtue of The Import of, and Trade in, Animals and Animal Products (Miscellaneous Amendments) (EU Exit) Regulations 2020 (S.I. 2020/1462), regs. 1(3), **51(11)** (with regs. 69-71)

ANNEX V U.K.

REPEALED DECISION WITH LIST OF ITS SUCCESSIVE AMENDMENTS

Commission Decision 2001/618/EC (OJ L 215, 9.8.2001, p. 48).	
Commission Decision 2001/746/EC (OJ L 278, 23.10.2001, p. 41).	Only as regards the reference to Decision 2001/618/EC in Article 1
Commission Decision 2001/905/EC (OJ L 335, 19.12.2001, p. 22).	Only as regards the reference to Decision 2001/618/EC in Article 2
Commission Decision 2002/270/EC (OJ L 93, 10.4.2002, p. 7).	Only Article 3
Commission Decision 2003/130/EC (OJ L 52, 27.2.2003, p. 9).	
Commission Decision 2003/575/EC (OJ L 196, 2.8.2003, p. 41).	
Commission Decision 2004/320/EC (OJ L 102, 7.4.2004, p. 75).	Only Article 2 and Annex II
Commission Decision 2005/768/EC (OJ L 290, 4.11.2005, p. 27).	
Commission Decision 2006/911/EC (OJ L 346, 9.12.2006, p. 41).	Only as regards the reference to Decision 2001/618/EC in Article 1 and point 12 of the Annex
Commission Decision 2007/603/EC (OJ L 236, 8.9.2007, p. 7).	
Commission Decision 2007/729/EC (OJ L 294, 13.11.2007, p. 26).	Only as regards the reference to Decision 2001/618/EC in Article 1 and point 10 of the Annex

ANNEX VI U.K.

CORRELATION TABLE

Decision 2001/618/EC	**This Decision**
Article 1(a) and (b)	Article 1, points 1 and 2

Article 1(c) first to fifth indent	Article 1, point 3(a) to (e)
Article 1(d) first to fourth indent	Article 1, point 4(a) to (d)
Article 2(a) and (b)	Article 2, points 1 and 2
Article 2(c) first to third indent	Article 2, point 3(a) to (c)
Article 3(a)	Article 3, point 1
Article 3(b) first and second indent	Article 3, point 2(a) and (b)
Article 3(c) first to sixth indent	Article 3, point 3(a) to (f)
Article 4(a)	Article 4, point 1
Article 4(b) first and second indent	Article 4, point 2(a) and (b)
Article 4(c) first to fifth indent	Article 4, point 3(a) to (e)
Articles 5 to 8	Articles 5 to 8
Article 9	—
Article 10	—
—	Article 9
Article 11	Article 10
Annexes I to IV	Annexes I to IV
—	Annex V
—	Annex VI

(**1**) OJ 121, 29.7.1964, p. 1977/64. Directive as last amended by Commission Decision 2007/729/EC (OJ L 294, 13.11.2007, p. 26).

(**2**) OJ L 215, 9.8.2001, p. 48. Decision as last amended by Decision 2007/729/EC.

(**3**) See Annex V.

Changes to legislation:
There are currently no known outstanding effects for the Commission Decision of 21 February 2008 on additional guarantees in intra-Community trade of pigs relating to Aujeszky's disease and criteria to provide information on this disease (notified under document number C(2008) 669) (Codified version) (Text with EEA relevance) (2008/185/EC).